AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A pharmaceutical composition for systemic administration comprising a pharmaceutically suitable carrier or diluent and a compound having the structure:

or pharmaceutically acceptable derivative thereof;

wherein R₁ is hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl;

R₂ and is methyl;

R₃ are each independently is hydrogen, halogen, hydroxyl, protected hydroxyl, or an aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl moiety; or

R₁-and-R₂, when taken together, may form a substituted-or-unsubstituted, saturated or unsaturated eyelic ring of 3 to 8 earbon atoms; or

R₁ and R₃, when taken together, may form a substituted or unsubstituted, saturated or unsaturated cyclic ring of 3 to 8 carbon atoms;

R4 is hydrogen or halogen;

R₅ is hydrogen, an exygen-protecting group or a protecting group;

R₆ is hydrogen, hydroxyl, or protected hydroxyl;

n is 0-2:

R₇, for each occurrence, is independently hydrogen, hydroxyl, or protected hydroxyl;

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R₈ is hydrogen, halogen, hydroxyl, protected hydroxyl, alkyloxy, or an aliphatic moiety optionally substituted with hydroxyl, protected hydroxyl, SR₁₂, or NR₁₂R₁₃;

 R_9 is hydrogen, halogen, <u>hydroxyl</u>, <u>protected hydroxyl</u>, OR_{12} , SR_{12} , $NR_{12}R_{13}$, - $X_1(CH_2)_pX_2-R_{14}$, or is lower alkyl optionally substituted with hydroxyl, protected hydroxyl, halogen, amino, protected amino, or $-X_1(CH_2)_pX_2-R_{14}$;

wherein R_{12} and R_{13} are, independently for each occurrence, hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl; or a protecting group, or R_{12} and R_{13} , taken together may form a saturated or unsaturated cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms, and each of R_{12} and R_{13} are optionally further substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen,

wherein X_1 and X_2 are each independently absent, or are oxygen, NH, or - N(alkyl), or wherein X_2 - R_{14} together are N_3 or are a saturated or unsaturated heterocyclic moiety,

p is 2-10, and

 R_{14} is hydrogen, or an aryl, heteroaryl, alkylaryl, or alkylheteroaryl moiety, or is – (C=O)NHR₁₅ –(C=O)OR₁₅, or –(C=O)R₁₅, wherein each occurrence of R_{15} is independently hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl; or R_{14} is –SO₂(R_{16}), wherein R_{16} is an aliphatic moiety, wherein one or more of R_{14} , R_{15} , or R_{16} are optionally substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen; or

R₈ and R₉ may, when taken together, form a saturated or unsaturated cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms and is optionally substituted with hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen;

R₁₀ is hydrogen, hydroxyl, protected hydroxyl, amino, or protected amino;

R₁₁ is hydrogen, hydroxyl or protected hydroxyl;

X is O, NH or CH2;

Y is CHR₁₇, O. CR₁₇ or NR₁₇; and Z is CHR₁₈, C=O, CR₁₈ or NR₁₈, wherein each occurrence of R_{17} and R_{18} is independently hydrogen or aliphatic, or R_{17} and R_{18} taken together

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is -O-, -CH2- or -NR19-, wherein R19 is hydrogen or lower alkyl, and Y and Z may be connected by a single or double bond;

wherein the compound is present in an amount effective to inhibit production of a proinflammatory and/or immunologic cytokine.

(Currently Amended) The composition of claim 1, wherein: 2.

R1 is hydrogen, straight or branched lower alkyl, straight or branched lower heteroalkyl, or aryl,

wherein the alkyl, heteroalkyl, and aryl groups may optionally be substituted with one or more occurrences of halogen, hydroxyl or protected hydroxyl;

R₂ and is methyl:

R₃ are each independently is hydrogen, halogen, hydroxyl, protected hydroxyl, straight or branched lower alkyl, straight or branched lower heteroalkyl, or aryl,

wherein the alkyl, heteroalkyl, and aryl groups may optionally be substituted with one or more occurrences of halogen, hydroxyl or protected hydroxyl; or

R4-and R2, when taken together, may form a saturated or unsaturated eyelic ring of 3 to 8 carbon atoms, optionally substituted with one or more occurrences of halogen; or

R₁ and R₃, when taken together, may form a saturated or unsaturated cyclic ring of 3 to 8 carbon atoms, optionally substituted with one or more occurrences of halogen;

R4 is hydrogen or halogen;

R₅ is hydrogen or a protecting group;

R₆ is hydrogen, hydroxyl, or protected hydroxyl;

n is 0-2:

R₁, for each occurrence, is independently hydrogen, hydroxyl, or protected hydroxyl;

R₈ is hydrogen, halogen, hydroxyl, protected hydroxyl, alkyloxy, or lower alkyl optionally substituted with hydroxyl, protected bydroxyl, SR₁₂, or NR₁₂R₁₃;

R₉ is hydrogen, halogen, hydroxyl, protected hydroxyl, OR₁₂, SR₁₂, NR₁₂R₁₃, - $X_1(CH_2)_pX_2-R_{14}$, or is lower alkyl optionally substituted with hydroxyl, protected hydroxyl, halogen, amino, protected amino, or -X₁(CH₂)_pX₂-R₁₄;

wherein R₁₂ and R₁₃ are, independently for each occurrence, hydrogen, lower alkyl, aryl, heteroaryl, alkylaryl, or alkylheteroaryl, or a protecting group, or R_{12} and R_{13} , taken together may form a saturated or unsaturated cyclic ring containing 1 to 4 carbon

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atoms and 1 to 3 nitrogen or oxygen atoms, and each of R_{12} and R_{13} are optionally further substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen,

wherein X1 and X2 are each independently absent, or are oxygen, NH, or -N(alkyl), or wherein X2-R14 together are N3 or are a saturated or unsaturated heterocyclic moiety,

p is 2-10, and

R₁₄ is hydrogen, or an aryl, heteroaryl, alkylaryl, or alkylheteroaryl moiety, or is -(C=O)NHR₁₅ -(C=O)OR₁₅, or -(C=O)R₁₅, wherein each occurrence of R_{15} is independently hydrogen, alkyl, heteroalkyl, aryl, heteroaryl, alkylaryl, or alkylheteroaryl, or R₁₄ is -SO₂(R₁₆), wherein R₁₆ is an alkyl moiety, wherein one or more of R₁₄, R₁₅, or R₁₆ are optionally substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen; or

R₈ and R₉ may, when taken together, form a saturated or unsaturated cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms and is optionally substituted with hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen;

R₁₀ is hydrogen, hydroxyl, protected hydroxyl, amino, or protected amino;

R11 is hydrogen, hydroxyl or protected hydroxyl;

X is 0;

Y is CHR₁₇, CR₁₇ or NR₁₇; and Z is CHR₁₈, C=O, CR₁₈ or NR₁₈, wherein each occurrence of R₁₇ and R₁₈ is independently hydrogen or lower alkyl, or R₁₇ and R₁₈ taken together is -O-, -CH2- or -NR19-, wherein R19 is hydrogen or lower alkyl, and Y and Z may be connected by a single or double bond.

- (Previously Presented) The composition of claim 2, where n is 1. 3.
- (Original) The composition of claim 2, where R₄ is halogen. 4.
- (Original) The composition of claim 2, where R₄ is fluorine. 5.
- (Original) The composition of claim 2, where Y and Z together represent-CH=CH-6.

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- 7. (Original) The composition of claim 2, where Y and Z together represent trans CH=CH-.
- 8. (Previously Presented) The composition of claim 2, wherein R_1 and R_2 are each methyl and R_3 is hydrogen and the compound has the structure:

wherein R₄-R₁₁, n, Y and Z are as defined in claim 2.

- 9. (Previously Presented) The composition of claim 8, wherein n is 1.
- 10. (Original) The composition of claim 8, wherein R₄ is halogen.
- 11. (Original) The composition of claim 8, wherein Y and Z together represent -CH=CH.
- 12. (Previously Presented) The composition of claim 8, wherein n is 1, R₄ is halogen and Y and Z together represent -CH=CH-.
- 13. (Original) The composition of claim 11 or 12 wherein -CH=CH- is trans.
- 14. (Previously Presented) The composition of claim 2, wherein R₉ is NR₁₂R₁₃ and the compound has the structure:

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wherein R₁-R₁₃, n, Y and Z are as defined in claim 2, or

R₁₃ and R₈ may, when taken together, form a cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms and is optionally substituted with hydrogen, alkyloxy, amino, alkylamino, aminoalkyl, and halogen.

- 15. (Previously Presented) The composition of claim 14, wherein n is 1.
- 16. (Original) The composition of claim 14, wherein R₄ is halogen.
- 17. (Original) The composition of claim 14, wherein Y and Z together represent -CH=CH-.
- 18. (Original) The composition of claim 14, wherein R_1 and R_2 are each methyl and R_3 is hydrogen.
- 19. (Previously Presented) The composition of claim 14, wherein n is 1, R₁ and R₂ are each methyl, R₃ is hydrogen, R₄ is halogen, and Y and Z together represent -CH=CH-.
- 20. (Original) The composition of claim 17 or 19, wherein -CH=CH- is trans.
- 21. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

22. (Original) The composition of claim 1 wherein the compound has the structure:

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23. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

24. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

25. (Previously Presented) The composition of claim 1 wherein the compound has the structure:

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(Previously Presented) The composition of claim 1 wherein the compound has the 26. structure:

or pharmaceutically acceptable derivative thereof.

(Original) The composition of claim 1 wherein the compound has the structure: 27.

or pharmaceutically acceptable derivative thereof.

(Original) The composition of claim 1 wherein the compound has the structure: 28.

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29. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

30. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

31. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

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32, (Previously Presented) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

33. (Original) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

34. (Previously Presented) The composition of claim 1 wherein the compound has the structure:

or pharmaceutically acceptable derivative thereof.

35. (Previously Presented) The composition of claim 1 wherein the compound has the structure:

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- 36. (Original) The pharmaceutical composition of claim 1, wherein the composition is for oral administration.
- 37. (Canceled)
- 38. (Previously Presented) The pharmaceutical composition of claim 1, wherein the proinflammatory and/or immunologic cytokine is TNFα, IL-1, IL-6, IL-8 or IL-2.
- 39. (Currently Amended) A method for treating rheumatoid arthritis, psoriasis, asthma, sepsis, inflammatory bowel disease, atopic dermatitis or Crohn's disease comprising the step of systemically administering to a subject in need thereof a pharmaceutically suitable carrier or diluent and a therapeutically effective amount of a compound having the structure: a therapeutically effective amount of a pharmaceutical composition of claim 1.

or pharmaceutically-acceptable derivative thereof;

wherein R₁ is hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl;

R₂-and R₃ are each independently hydrogen, halogen, hydroxyl, protected hydroxyl, or an aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl moiety; or

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R₁-and R₂, when taken together, may form a substituted or unsubstituted, saturated or unsubstituted, saturated

R₁-and R₂, when taken together, may form a substituted or unsubstituted, saturated or unsubstituted, saturated

R4 is hydrogen or halogen;

Rs is hydrogen, an exygen protecting group or a prodrug;

R6 is hydrogen, hydroxyl, or protected hydroxyl;

n is 0 2;

R₇, for each occurrence, is independently hydrogen, hydroxyl, or protected hydroxyl;

R₈ is-hydrogen, halogen, hydroxyl, protected hydroxyl, alkyloxy, or an aliphatic moiety optionally substituted with hydroxyl, protected hydroxyl, SR₁₂, or NR₁₂R₁₂;

R₉ is hydrogen, halogen, hydroxyl, protected hydroxyl, OR₁₂, SR₁₂, NR₁₂R₁₃, X₁(CH₂)_pX₂-R₁₄, or is lower alkyl optionally substituted with hydroxyl, protected hydroxyl, halogen, amino, protected amino, or -X₁(CH₂)_pX₂-R₁₄;

wherein R₁₂ and R₁₃ are, independently for each occurrence, hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl; or a protecting group, or R₁₂ and R₁₃, taken together may form a saturated or unsaturated cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms, and each of R₁₂ and R₁₃ are optionally further substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen,

wherein X₁ and X₂ are each-independently absent, or are exygen, NH, or N(alkyl), or wherein X₂ R₁₄ together are N₂ or are a saturated or unsaturated heterocyclic moiety,

p is 2 10, and

R₁₄ is hydrogen, or an aryl, heteroaryl, alkylaryl, or alkylheteroaryl moiety, or is (C=O)NHR₁₅ (C=O)OR₁₅, or (C=O)R₁₅, wherein each occurrence of R₁₅—is independently hydrogen, aliphatic, heteroaliphatic, alicyclic, heteroalicyclic, aryl or heteroaryl; or R₁₄ is SO₂(R₁₆), wherein R₁₆ is an aliphatic moiety, wherein one or more of R₁₄, R₁₅, or R₁₆ are optionally substituted with one or more occurrences of hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen; or

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Rs and Ro may, when taken together, form a saturated or unsaturated eyelic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or exygen atoms and is optionally substituted with hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen;

R₁₀-is hydrogen, hydroxyl, protected hydroxyl, amino, or protected amino;

R11 is hydrogen, hydroxyl or protected hydroxyl;

Y is CHR17, CR17 or NR17; and Z is CHR18, C-O, CR18 or NR18, wherein each occurrence of R₁₇ and R₁₈ is independently hydrogen or aliphatic, or R₁₇ and R₁₈ taken together is O, CH2 or NR19, wherein R19 is hydrogen or lower alkyl, and Y and Z may be connected by a single or double bond;

wherein the compound is present in an amount effective to inhibit production of a proinflammatory and/or immunologic cytokine.

- 40. (Original) The method of claim 39, wherein the compound is administered orally.
- 41. (Canceled)
- 42. (Previously Presented) The method of claim 39, wherein the method is for treating psoriasis.
- (Currently Amended) The method of claim 41, wherein the compound has any one of 43. the following structures:

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44. (Canceled)

- 45. (Previously Presented) The method of claim 39, wherein the pro-inflammatory and/or immunologic cytokine is TNFα, IL-1, IL-6, IL-8 or IL-2.
- 46. (New) The composition of claim 2, where R₁ is hydrogen or methyl.
- 47. (New) The composition of claim 2, where R₃ is hydrogen or halogen.
- 48. (New) The composition of claim 2, where R₄ is hydrogen.
- 49. (New) The composition of claim 2, where R₅ is hydrogen.
- 50. (New) The composition of claim 2, where R₆ is hydroxyl.
- 51. (New) The composition of claim 2, where R₇ is hydrogen or hydroxyl.
- 52. (New) The composition of claim 2, where R₈ is hydrogen or halogen.

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- 53. (New) The composition of claim 2, where R_9 is hydroxyl, protected hydroxyl, $-OR_{12}$, $-NR_{12}R_{13}$ or $-O(CH_2)_pX_2R_{14}$, wherein R_{12} , R_{13} , R_{14} and X_2 are as defined in claim 2.
- 55. (New) The composition of claim 53, where R_9 is -NR₁₂R₁₃, wherein R₁₂ is methyl, ethyl, propyl, isopropyl, or butyl, optionally substituted with one or more occurrences of hydroxyl or protected hydroxyl, and R₁₃ is hydrogen or lower alkyl, or NR₁₂R₁₃ together represents a 5- or 6-membered heterocyclic moiety.
- 56. (New) The composition of claim 53, where R₉ is -O(CH₂)_pX₂R₁₄, wherein X₂R₁₄ together represent N₃, NMe₂, NHAc, NHSO₂Me, NHCONHMe, NHCONHPh, morpholine, imidazole, aminopyridine, or any one of:

- 57. (New) The composition of claim 2, where R₈ and R₉, taken together, form a saturated or unsaturated cyclic ring containing 1 to 4 carbon atoms and 1 to 3 nitrogen or oxygen atoms and is optionally substituted with hydroxyl, protected hydroxyl, alkyloxy, amino, protected amino, alkylamino, aminoalkyl, or halogen.
- 58. (New) The composition of claim 2, where R_{10} is hydroxyl.
- 59. (New) The composition of claim 2, where R₁₁ is hydrogen.
- 60. (New) The composition of claim 2, where Y and Z together are cyclopropyl.

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61. (New) The composition of claim 2, where Y and Z together are -NHC(=O)-.